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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	NO. CONFIRMATION NO.	
10/748,939	12/30/2003	Markus A. Wicki	59457US002	1916	
32692 3M INNOVAT	7590 07/05/2007 FIVE PROPERTIES COM	EXAMINER			
PO BOX 3342	7	OH, TAYLOR V			
ST. PAUL, M	N 33133-3427		ART UNIT PAPER NUMBER		
		1625			
			NOTIFICATION DATE	DELIVERY MODE	
			07/05/2007	ELECTRONIC	

Please find below and/or attached an Office communication concerning this application or proceeding.

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	Application No.	Applicant(s)
	10/748,939	WICKI ET AL.
Office Action Summary	Examiner	Art Unit
•	Taylor Victor Oh	1625
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA  - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailling date of this communication.  - If NO period for reply is specified above, the maximum statutory period w  - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be timulated and will expire SIX (6) MONTHS from a cause the application to become ABANDONEI	l. ely filed the mailing date of this communication. O (35 U.S.C. § 133).
Status		
<ul> <li>1) Responsive to communication(s) filed on 06 Ju</li> <li>2a) This action is FINAL. 2b) This</li> <li>3) Since this application is in condition for allowar closed in accordance with the practice under E</li> </ul>	action is non-final. nce except for formal matters, pro	
Disposition of Claims		
4) ☐ Claim(s) 1,2,4,5,7,8,10,11,13-17 and 20-30 is/a 4a) Of the above claim(s) is/are withdray 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) 1-2,4-5,7-8,10-11,13-17,20-30 is/are r 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) are subject to restriction and/or	vn from consideration. rejected.	
Application Papers		
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) access applicant may not request that any objection to the Replacement drawing sheet(s) including the correct and the other controls.  The oath or declaration is objected to by the Examine	epted or b) objected to by the Eddrawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		•
<ul> <li>12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:</li> <li>1. Certified copies of the priority documents</li> <li>2. Certified copies of the priority documents</li> <li>3. Copies of the certified copies of the priority application from the International Bureau</li> <li>* See the attached detailed Office action for a list of the certified copies of the priority application from the International Bureau</li> </ul>	s have been received. s have been received in Application rity documents have been receive u (PCT Rule 17.2(a)).	on No ed in this National Stage
Attachmant/al		
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO/SB/08)  Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	ite

## Continued Examination Under 37 CFR 1.114

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 6/06/07 has been entered.

## The Status of Claims

Claims 1-2,4-5,7-8, 10-11,13-17, 20-30 are pending.

Claims 1-2,4-5,7-8, 10-11,13-17, 20-30 are rejected.

#### **DETAILED ACTION**

1. Claims 1-2,4-5,7-8, 10-11,13-17, 20-30 are under consideration in this Office Action.

### **Priority**

2. None.

# **Drawings**

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3. None.

# Claim Rejections - 35 USC § 103

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

The factual inquiries set forth in *Graham* v. *John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

- 1. Determining the scope and contents of the prior art.
- 2. Ascertaining the differences between the prior art and the claims at issue.
- 3. Resolving the level of ordinary skill in the pertinent art.

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4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

Claims 1-2,4-5,7-8, 10-11,13-17, 20-30 are rejected under 35 U.S.C. 103(a) as being unpatentable over Babler et al (Tetrahedron Letters, 1979, no. 22, p. 1971-74).

Babler et al teaches a selective esterification method of treating diol with a solution of acetic acid in the presence of sulfuric acid at a room temperature to produce the monoacetate (see page 1971, lines 17-19)free from the corresponding diacetate derivatives, which is done by continuous extraction with a suitable nonpolar solvent (see page 1973, lines 1-3). Furthermore, various starting diols compared with glacial acetic acid are described below (see page 1972, table 1):

		1.	ABLE I			
	Aqueous Reaction			Product Elstributions		
	Histore (al of H <sub>2</sub> D: ml of glacial acetic acid: ml of conc. H <sub>2</sub> SO <sub>4</sub> )	Tineb	Solvent Used for Extraction <sup>C</sup>	Diol	(1) Mondécetéte <sup>©</sup>	Discetate
15 resol of 1,10-decane- diol	155:75:0.25	36 hrs	5:1 (v/v) cyclohexane: CC14	39 <sup>‡</sup>	60	1
is resol of 1,10-decame- dial	120:50:0.25	35 hrs	l:l (v/v) hexane: cyclohe:Ane	824	75	3
IS resol of ethylene glycol	160:80:0.50	1 wk	bonzene	0	94	6
17 mol of 1.4-cyclohex- enedio19	180:45:4.0	4 days	benzene	14h	85 5	1
7 mmpl of 1,12-dodecane 3501	90:150:n.25	30 hrs	cycloheza <del>za</del>	29 <sup>‡</sup>	66	5
13.7 mmol of 1,8-actane- Hol	200:25:4.0	40 hrs	hexane	2	94	4

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acid. This mixture was subsequently extracted continuously with the specified nonpolar solvent. Differ time represents that required for essentially quantitative removal of the starting diol (most of which has been converted to the corresponding monoacetate) from the aqueous reaction mixture using the specific reaction conditions listed in the table. This time should be able to be reduced substantially by increasing the amount of sulfuric acid catalyst reating the aqueous reaction mixture to a moderate temperature (e.g., 50°C), varying the ratio of water-acetic acid (up to a certain point), and using a more efficient extractor. The solvents utilized in the above reactions do not necessarily represent the optimum one for each particular system. The latter can be ascertained only after extensive development of this process. Suitable nonpolar solvents include alkanes, cycloalkanes, aromatic hydrocarbons, halide derivatives of hydrocarbons, or mixtures thereof. The product mixture was isolated by cooling the solvent used for extraction to room temperature. Any unreacted diol present in the product mixture often precipitated out of the nonpolar solvent at this stage and could be recovered by simple filtration. After drying the filtrate over anhydrous K2CO3 and subsequent removal of the extraction solvent under reduced pressure, product ratios were determined by VPC analysis (6'x1/8" SE-30 column). Retention times: diester > manoacetate

(see page 1972, a lower part of paragraph).

To illustrate the utility of this selective esterification process, 1.8-octanediol monoacetate<sup>8</sup> (2. n=6) was converted in two steps to the sex pheromone of the oriental fruit moth,<sup>9</sup> a pest of peach orchards. Oxidation of the unprotected hydroxyl group using pyridinium chlorochromate<sup>10</sup> afforded the previously reported<sup>11</sup> 8-acetoxyoctanal in 86% yield. Subsequent treatment of the latter aldehyda with the yild derived from n-butyl-triphenylphosphonium bromide<sup>12</sup> under "salt-free" conditions<sup>13</sup> gave Z-8-dodecenyl acetate,<sup>14</sup> > 98% pure by YPC analysis, <sup>15</sup> in 45% yield.

(see

page 1973, a middle paragraph).

The instant invention, however, differs from the prior art in that the carboxylic acid has a solubility in water of at least 20 % by weight at 20 °C; the aprotic solvent has a polarity index between 1.5 and 3.5; the reaction step is conducted as a batch process.

With respect to the solubility of the acetic acid in water, it is well-known that the acetic acid is very soluble in water. Therefore, it would have been obvious to the skilled artisan in the art to be motivated to find the solubility of the acetic acid in water by routine experimentation in order to discover the optimum range of the acetic acid in water for the reaction process.

Concerning the polarity index of the aprotic solvent, the reference is silent about it. However, the limitation of , pH, concentration , a polarity index does not impart patentability to a process when such values are those which would be determined by one of ordinary skill in the art in achieving optimum operation of the process. The polarity index of the solvent is well understood by those of ordinary skill in the art to be a result-effective variable, especially when attempting to control selectivity of a chemical process.

With respect to the claimed reaction step being conducted as a batch process, the prior art process is continuous as in the case of the extraction step. However, according to the a case law, it is well- established that batch and continuous processes are not patentably distinct. See , e.g. , In re Dilnot, 319 F. 2d 188, 138 USPQ 248 (C.C.P. A. 1963). Therefore, it would have been obvious to the skilled artisan in the art to be motivate to conduct the prior art process as a batch process as an alternative.

Babler et al expressly teaches the selective esterification method of treating diol with a solution of acetic acid in the presence of sulfuric acid at a room temperature to produce the pure monoacetate by continuous extraction with the suitable nonpolar solvent (see page 1973, lines 1-3). The good selection of the polarity index of the solvent can be useful in attempting to control selectivity of a chemical process. Therefore, it would have been obvious to the skilled artisan in the art to be motivated to optimize the continuous extraction with the suitable nonpolar solvent by routine

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experimentation on the selection of the polarity index of the solvent in order to maximize

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the desired product for the reaction process.

Any inquiry concerning this communication or earlier communications from the

examiner should be directed to Taylor Victor Oh whose telephone number is 571-272-

0689. The examiner can normally be reached on 8:30-5:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Janet Andres can be reached on 571-272-0867. The fax phone number for

the organization where this application or proceeding is assigned is 571-273-8300.

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USPTO Customer Service Representative or access to the automated information

system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Taylor Victor Oh, MSD,LAC

6/21/01

**Primary Examiner** 

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